to an amino acid sequence of SEQ ID NO:1,

- c) a biologically active fragment of an amino acid sequence of SEQ ID NO:1, and
- d) an immunogenic fragment of an amino acid sequence of SEQ ID NO:1.



- 11. (Twice Amended.) A [pharmaceutical] composition comprising an effective amount of a polypeptide of claim 1 in conjunction with a suitable pharmaceutical carrier.
- 12. (Reiterated.) A purified antibody which binds specifically to a polypeptide of claim 1.
- 13. (Reiterated.) A purified agonist which specifically binds to and modulates the activity of a polypeptide of claim 1.
- **14.** (**Reiterated.**) A purified antagonist which specifically binds to and modulates the activity of a polypeptide of claim 1.
- **15.** (**Reiterated.**) A pharmaceutical composition comprising an antagonist of claim 14 in conjunction with a suitable pharmaceutical carrier.
- 16. (Reiterated.) A method for treating liver disease comprising administering to a subject in need of such treatment an effective amount of a pharmaceutical composition of claim 15.
- 17. (Reiterated.) A method for detection of a polynucleotide encoding a polypeptide comprising SEQ ID NO:1 in a biological sample, said method comprising the steps of:
- a) hybridizing an isolated and purified polynucleotide which is complementary to a polynucleotide encoding a polypeptide comprising SEQ ID NO:1 to nucleic acid material of a biological sample; and
- b) detecting said hybridization complex, wherein the presence of said complex correlates with the presence of a polynucleotide encoding a polypeptide comprising SEQ ID NO:1 in said biological sample.

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